"Lee's Milk of Magnesia Dental Cream * * The Trade Laboratories, Inc.,

Distributors, Newark, N. J."

The article was alleged to be misbranded in that the representation in the labeling that it would make the gums healthy and firm, was false and misleading, since it was not efficacious for the purposes recommended.

It was also alleged to be misbranded under the provisions of the law ap-

plicable to cosmetics reported in C. N. J. No. 24.

On September 25, 1939, no claimant having appeared, judgment of condemnation was entered and the product was ordered delivered to a charitable organization.

MISCELLANEOUS

74. Adulteration and misbranding of epinephrine chloride solution. U. S. v. One Carton of Solution Epinephrine Chloride. Default decree of condemnation and destruction. (F. D. C. No. 596. Sample No. 51848-D.)

This product had a potency of not more than 65 percent of the requirement of

the United States Pharmacopoeia for epinephrine hydrochloride.

On September 14, 1939, the United States attorney for the District of New Jersey filed a libel against one carton, containing seven 1-ounce bottles, of epinephrine chloride solution at Camden, N. J., alleging that the article had been shipped in interstate commerce on or about April 29, 1939, by Harvey-Pittenger Co. from Philadelphia, Pa.; and charging that it was adulterated and misbranded.

It was alleged to be adulterated in that it purported to be or was represented as a drug the name of which is recognized in the United States Pharmacopoeia, and its strength differed from or its quality fell below the standard set forth in the pharmacopoeia.

It was alleged to be misbranded in that the representation in the labeling that each fluid ounce contained 0.45 grain of epinephrine was false and misleading since it contained less than 0.45 grain of epinephrine in each fluid

ounce.

On October 20, 1939, no claimant having appeared, judgment of condemnation was entered and the product was ordered destroyed.

75. Adulteration and misbranding of Daily Vitamins. U. S. v. 84 Boxes of Daily Vitamins. Default decree of condemnation and destruction. (F. D. C. No. 556. Sample No. 47472–D.)

This product was represented to contain 200 International Units of vitamin B₁ per capsule, whereas it contained not more than 10 International Units of

vitamin B1 per capsule.

On September 5, 1939, the United States attorney for the District of Columbia filed a libel against 84 boxes of Daily Vitamins at Washington, D. C., alleging that the article had been shipped in interstate commerce on or about December 22, 1938, by Daily Vitamins, Inc., from Cincinnati, Ohio; and charging that it was adulterated and misbranded.

It was alleged to be adulterated in that its strength differed from and its purity fell below that which it purported or was represented to possess.

It was alleged to be misbranded in that the representation in the labeling that each capsule contained not less than 400 Sherman Units (200 International Units) of vitamin B₁, was false and misleading.

On September 26, 1939, no claimant having appeared, judgment of condem-

nation was entered and the product was ordered destroyed.

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Issued May 1940

United States Department of Agriculture

After

FOOD AND DRUG ADMINISTRATION

NOTICES OF JUDGMENT UNDER THE FEDERAL FOOD, DRUG, AND COSMETIC ACT

[Given pursuant to section 705 of the Food, Drug, and Cosmetic Act]

76-140

DRUGS AND DEVICES

The cases reported herein were instituted in the United States district courts by the United States attorneys, acting upon reports submitted by direction of the Secretary of Agriculture.

GROVER B. HILL, Acting Secretary of Agriculture.

Washington, D. C., May 8, 1940.

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MISBRANDED SEDATIVES, PAIN RELIEVERS, AND HEADACHE REMEDIES

76. Misbranding of Causalin. U. S. v. Amfre Drug Co., Inc., and Lewis Stern. Pleas of guilty. Fines, \$1,400. (F. D. C. No. 94. Sample Nos. 25962-D, 25963-D, 25964-D, 30071-D to 30074-D, incl., 30092-D to 30097-D, incl., 35452-D, 35453-D, 35567-D, 35569-D, 35570-D, 41997-D.)

This product consisted of capsules and tablets containing aminopyrine (aminodimethylpyrazolon,) salicylic ethyl ester carbonate, and a sulfonate such as quinolinesulfonate. It would be dangerous to health when used in the dosage or with the frequency prescribed, recommended, and suggested in the labeling.

The labeling of 6 of the 7 shipments bore the recommendation that the article be taken in the dosage as directed by the physician, 1 to 2 tablets or capsules 3 times a day, one-half hour before meals. In the seventh shipment the labeling in describing the "manner of use" of the article represented that the average dose was 1 capsule or tablet 3 times daily one-half hour before meals and that in severe or chronic cases one should start with 2 capsules or tablets 3 times daily continuing for about a week, then gradually reducing the dosage.

On January 30, 1940, the United States attorney for the Southern District of New York filed an information against the Amfre Drug Co., Inc., and Lewis Stern, president of the corporation, alleging shipment by said defendants within the period from on or about July 1 to on or about December 27, 1938, from the State of New York into the States of New Jersey, Pennsylvania, Massachusetts, and Rhode Island, of quantities of Causalin which was misbranded for the reasons appearing above.